Assessment of Unabated Facility
Emission Potentials for Evaluating
Airborne Radionuclide Monitoring
Requirements at Pacific Northwest
National Laboratory – 1999

September 1999

Prepared for the U.S. Department of Energy under Contract DE-AC06-76RLO 1830

Pacific Northwest National Laboratory Richland, Washington 99352

DISCLAIMER

This report was prepared as an account of work sponsored by an agency of the United States Government. Neither the United States Government nor any agency thereof, nor any of their employees, make any warranty, express or implied, or assumes any legal liability or responsibility for the accuracy, completeness, or usefulness of any information, apparatus, product, or process disclosed, or represents that its use would not infringe privately owned rights. Reference herein to any specific commercial product, process, or service by trade name, trademark, manufacturer, or otherwise does not necessarily constitute or imply its endorsement, recommendation, or favoring by the United States Government or any agency thereof. The views and opinions of authors expressed herein do not necessarily state or reflect those of the United States Government or any agency thereof.

DISCLAIMER

Portions of this document may be illegible in electronic image products. Images are produced from the best available original document.

Summary

Assessments were performed to evaluate compliance with the airborne radionuclide emission monitoring requirements in the National Emission Standards for Hazardous Air Pollutants (NESHAP - U.S. Code of Federal Regulations, Title 40 Part 61, Subpart H). In these assessments, potential unabated offsite doses were evaluated for emission locations at facilities owned by the U.S. Department of Energy and operated by Pacific Northwest National Laboratory (Pacific Northwest) on the Hanford Site. Two of the facilities evaluated, 325 Building Radiochemical Processing Laboratory, and 331 Building Life Sciences Laboratory met state and federal criteria for continuous sampling of airborne radionuclide emissions. One other building, the 3720 Environmental Sciences Laboratory, was recognized as being in transition with the potential for meeting the continuous sampling criteria.

The assessments were performed using building radionuclide inventory data obtained in 1999. The list of buildings evaluated is provided in Table S.1.

Table S.1. PNNL-Operated DOE Buildings with Radioactive Inventories in 1999.

200 Environmental Monitoring Shed	331-G Interim Tissue Repository
2718E Critical Mass Fissile Storage	331-H Aerosol Wind Tunnel Research Facility
300-N Caisson	3718A Lab Equipment Central Pool
305-B Hazardous Waste Storage Facility	3720 Environmental Sciences Laboratory
306-W Materials Development Building	3730 Gamma Irradiation Facility
314 Engineering Development Laboratory	3745 Radiological Calibrations & Standards
	Building
318 Radiological Calibrations Laboratory	3020 Environmental Molecular Sciences
•	Laboratory
320 Analytical and Nuclear Research	622-R Meteorological Laboratory
Laboratory	
323 Mechanical Properties Laboratory	6652 Arid Land Ecology Laboratory
325 Radiochemical Processing Laboratory	747A Whole Body Counter
326 Materials Sciences Laboratory	Field Lysimeter Test Facility (FLTF)
329 Chemical Sciences Laboratory	Prototype Surface Barrier (PSB)
331 Life Sciences Laboratory	

Contents

Summaryii	ii
1.0 Introduction	
2.0 Assessment Methodology	.1
2.1 Projections of Annual Emission Quantities	1
2.2 Maximum Receptor Unit Dose Calculation	.4
2.3 Potential Emission Dose Assessment	.5
3.0 References	.1
Appendix A - Common Radionuclide Mixtures	.1
Appendix B - Radionuclide Inventory Database FeaturesB.	.1
Appendix C - Unit Dose Factor Calculations	.1
Tables	
S.1 PNNL-Operated DOE Buildings with Radioactive Inventories in 1999 i	iii
1 Common Radionuclide Mixtures	.3
2 Physical Forms and Potential Annual Release Fractions for Radionuclides	.4
3 Unit Dose Factors for 300 Area Source Location	.6
4 Emission System Potential Dose Assessment Summary	.8

1.0 Introduction

Requirements for sampling for airborne radionuclide emissions are contained in the following regulations and guides:

- U.S. Code of Federal Regulations (CFR), Title 40, Subpart H: National Emission Standards for Hazardous Air Pollutants (NESHAP)(1989)
- Washington Administrative Code (WAC) 246-247: Radiation Protection Air Emissions (1994)
- U.S. Department of Energy, DOE 5400.1: General Environmental Protection Program (1990)
- U.S. Department of Energy, DOE/EH-0173T, Environmental Regulatory Guide for Radiological Effluent Monitoring and Environmental Surveillance (January 1991).

These documents require that continuous sampling be performed at emission points that have the potential to cause an offsite dose of 0.1 mrem/yr if routine emissions were not mitigated by engineered pollution control systems. In addition, DOE 5400.1 specifies that a written plan be prepared for each facility that uses, generates, releases, or manages significant pollutants or hazardous materials. Thus, a Facility Effluent Monitoring Plan (FEMP) is prepared for facilities that require continuous sampling according to NESHAP.

In response to these requirements, the potential unmitigated offsite receptor dose from the facilities operated by the U.S. Department of Energy's (DOE's) Pacific Northwest National Laboratory^(a) (Pacific Northwest) containing radioactive materials or sources is evaluated annually. These evaluations were performed initially in 1991 for DOE's Pacific Northwest facilities on the Hanford Site. Based on the initial assessments, four Pacific Northwest buildings were identified as containing a sufficient inventory of radioactive material such that unmitigated emissions could potentially result in an annual offsite maximum receptor dose of 0.1 mrem. These buildings were the 324 Waste Technology Engineering Laboratory, the 325 Applied Chemistry Laboratory, the 327 Postirradiation Testing Laboratory, and the 3720 Chemistry and Material Sciences Laboratory. In accordance with the NESHAP, qualifying emission points from these buildings were sampled continuously. Also, in accordance with DOE 5400.1, FEMPs were prepared for these buildings.^(b)

The original radionuclide assessments were updated annually. There were no changes in the facility emission NESHAP status until 1996 when the 324 and 327 Buildings were transitioned to Babcock & Wilcox. In 1997, proposed movement of the radionuclide inventory from the 3720

⁽a) Operated by Battelle for the U.S. Department of Energy under contract DE-ACO6-76RLO 1830.

⁽b) The FEMPs have been updated over the years and are currently available for the 325, 331, 3720, and Balance-of-Plant Facilities (Pacific Northwest National Laboratory 199a,b,c,d).

Building to the 331 Building upgraded the 331 Building to major stack status. The move was accomplished in 1998 and 1999 with disposal of some materials resulting in both the 3720 and the 331 Buildings Potential-To-Emit¹ (PTE) dropping to less than 0.1 mrem. The major stack for 3720 may be downgraded to minor, but continued work in the 331 Building may result in increased throughput. Thus, the 331 Building main stack is maintained as a major stack and continuously sampled. Results of the 1992-1993 assessment were documented by Sula and Jette (1994), and results of the 1995 assessment were documented in Ballinger, Jette, and Sula (1995). This report documents the most recent methods and assessments for 1999.

¹ Potential-to-Emit is defined as the rate of release of radionuclides from an emission unit based on the actual or potential discharge of the effluent stream that would result if all abatement control equipment did not exist, but operations are otherwise normal.

2.0 Assessment Methodology

Requirements for facility air emission sampling are promulgated by

- the U.S. Environmental Protection Agency (EPA) in 40 CFR 61, "National Emissions Standards for Hazardous Air Pollutants," Subpart H, "National Emission Standards for Emissions of Radionuclides Other than Radon from Department of Energy Facilities"
- the Washington State Department of Health in Washington Administrative Code 246-247, "Regulation of Radioactive Air Emissions"
- the DOE in regulatory guide DOE/EH-0173T, "Environmental Regulatory Guide for Radiological Monitoring and Environmental Surveillance."

These regulations require that all emission units with the potential to emit radioactivity be evaluated, and that the emission units be continuously sampled if there is the potential for unmitigated releases to cause a dose of 0.1 mrem/yr to a maximum offsite receptor.

2.1 Projections of Annual Emission Quantities

Several methods for projecting potential unmitigated annual emission quantities are prescribed in the 1994 issuance of WAC 246-247:

- apply an annual release fraction to the radionuclide inventory in the facility
- multiply actual measured annual emissions by control system decontamination factors
- add actual measured annual emission quantities to actual measured quantities retained by control systems
- measure the annual discharge upstream from all control devices.

The inventory-based assessment method^(a) has been used by Pacific Northwest since the initial facility assessment in 1991. Whereas the inventory method yields an assessment based on the current building status (or even the future status if projected future inventory quantities are used in the assessment), the other prescribed methods yield an assessment based on past facility

⁽a) This method is described in WAC 246-247 as follows: Multiply the annual possession quantity of each radionuclide by the release fraction for that radionuclide, depending on its physical state. Use the following release fractions: (i) 1 for gases, (ii) 10⁻³ for liquids or particulate solids, and (iii) 10⁻⁶ for solids. Determine the physical state for each radionuclide by considering its chemical form and the highest temperature to which it is subjected. Use a release fraction of 1 if the radionuclide is subjected to temperatures at or above its boiling point; use a release fraction of 10⁻³ if the radionuclide is subjected to temperatures at or above its melting point but below its boiling point. If the chemical form is not known, use a release fraction of 1 for any radionuclide that is heated to a temperature of one hundred degrees Celsius or more, boils at a temperature of one hundred degrees Celsius or less, or is intentionally dispersed into the environment.

measurements. Thus, the inventory method may be more appropriate for use at research and development facilities where types and quantities of radionuclides may change from year to year and where historical sampling data may not be a reliable predictor of future emissions.

At Pacific Northwest, radioactive source and material information is maintained using three separate inventory systems:

Nuclear Materials Inventory. This inventory includes the majority of tritium, uranium, and transuranics in Pacific Northwest facilities. The inventory includes material in process as well as residual contamination from historical operations in the facility. The Nuclear Materials Inventory is categorized as "Type 3 inventory."

Composite Radioactive Materials Inventory. This inventory includes DOE-owned sealed radioactive sources above specific de minimus values (based on the December 1998 revision to 10 CFR 835) and all radioactive material that is possessed by Pacific Northwest under a State of Washington Radioactive Materials License. Most of the radionuclides in this inventory are in "sealed source" form. The Composite Radioactive Materials Inventory is categorized as "Type 2 inventory."

Facilities Management Radioactive Materials Inventory. This database was developed specifically to account for all radioactive material not included in either the Type 2 or Type 3 inventories. This category consists primarily of fission product radionuclides, including radionuclides in process as well as residual contamination (e.g., in hot cells) from historical operations in the facility. The Facilities Management Radioactive Material Inventory is categorized as "Type 1 inventory."

The Pacific Northwest Effluent Management Group annually requests Type 1 inventory information from the custodians of radioactive inventory. This request is made through a central point of contact for each of Pacific Northwest's Divisions who then request inventory information from custodians in their organization. Effluent Management also requests Type 2 and Type 3 inventories from the Composite Radioactive Materials Inventory and Nuclear Materials Inventory databases, respectively.

Although the type of information contained in each of the systems varies, after compiling the information and reformatting it, radioactive materials are identified by type, physical form, and quantity. Quantities are expressed either in terms of activity (Ci) or mass (grams). Radioactive material types may be expressed as specific radionuclides or as standard mixtures of radionuclides. For example, mixed radionuclides may be expressed as natural uranium, depleted uranium, Hanford 6%(²⁴⁰Pu) plutonium, etc., as shown in Table 1. Appendix A provides the basis for these values. In addition to the mixtures shown in the table, mixtures of fission products are assumed to consist of ⁹⁰Sr. Individual radionuclides or mixtures of radionuclides in small quantities may also be reported as *alpha* or *beta activity* for simplification. These materials are assumed to be ²⁴¹Am (alpha) or ⁹⁰Sr (beta) for dose assessment purposes.

Table 1. Common Radionuclide Mixtures

Material	Assumed Radionuclide	Specific Activity (Ci/g)
Depleted Uranium	²³⁸ U	3.64E-07
Natural Uranium	²³⁸ U	6.87E-07
U-enriched (<20% ²³⁵ U)	²³⁵ U	9.36E-06
U-enriched (<90% ²³⁵ U)	²³⁵ U	6.21E-05
Pu (6% ²⁴⁰ Pu)	²³⁹ Pu	8.00E-02
Pu (12% ²⁴⁰ Pu)	²³⁹ Pu	9.60E-02
Pu (24% ²⁴⁰ Pu)	²³⁹ Pu	3.48E-01

Radionuclides meeting any of the following criteria are excluded from the assessments:

- radionuclides present in commercially available building/construction materials
- radionuclides that can be purchased or possessed without a special radioactive materials license
- radionuclides <100 pCi/g alpha activity and <400 pCi/g beta activity.

Inventory data is entered into an ACCESS database that was developed specifically for these calculations. Appendix B provides a summary of the database features. The data are reviewed and revised as needed to eliminate duplicates, provide consistency in nuclide and unit identifications, and obtain additional information as required by the calculations. The review process is documented and filed with the assessment records.

Potential release fractions for radionuclides are based on the physical form of the radionuclide as shown in Table 2. Radionuclides present as sealed sources or in sealed, unvented Department of Transportation (DOT) shipping containers are assumed to be unavailable for release under normal circumstances.

Table 2. Physical Forms and Potential Annual Release Fractions for Radionuclides

Form	Code	Description	Potential Release Fraction
Gas	G	Nuclide will exceed its boiling point when uncontained, except that nuclides in gaseous form in commercial gas cylinders that are not opened may be listed as L.	1
Liquid/Powder	L/P	Nuclide will exceed its melting point or be present in particulate form (Aerodynamic Mean Diameter [AMD] <100 microns) when uncontained, except liquid and powders in unopened containers may be listed as S.	10 ⁻³
Solid	S	Nuclides not meeting conditions for the more dispersible classes.	10-6
Contained	С	Sealed sources or material in sealed, DOT containers, except those meeting exempt criteria.	0
Exempt	E	Sealed sources engineered to pass the special form testing specified by the DOT in 40CFR173.469 or ANSI ^(a) N43.6, or sealed in Type B DOT shipping containers.	0

⁽a) American National Standards Institute.

2.2 Maximum Receptor Unit Dose Calculation

For the unit dose calculations, the maximum offsite receptor is defined as an individual whose residence location, work location, and lifestyle maximize the dose from airborne pathways. All potential environmental transport pathways associated with an airborne radionuclide release were included (i.e., air inhalation, air submersion, exposure to deposited radionuclides, uptake of vegetation grown in contaminated soil). In addition, site-specific atmospheric dispersion and environmental transport and uptake parameters were used (see Appendix C).

Unit dose factors for the maximum offsite receptor were calculated for specific radionuclides using the EPA compliance code CAP88-PC (Parks 1992). For radionuclides that were not represented in CAP88-PC, GENII (Napier et al. 1988) was used to calculate unit doses. Radionuclides that were not represented in either CAP88-PC or GENII were assigned default values equal to that of ²⁴¹Am for alpha emitters or ⁹⁰Sr for non-alpha emitters. The decay of the daughter products was also considered for short half-life radionuclides in assigning default

values. The unit dose factor calculations were performed for a single source point in the 300 Area of the Hanford Site (Table 3) and a single source point in the 200 East Area of the Site (WHC 1991).

For Building 6652 (located in the Fitzner/Eberhardt Arid Land Ecology [ALE] Reserve), dose assessments were performed by applying a *location correction factor* to the 300 Area unit dose factor to correct for varying source-receptor distances and directions. The location correction factor was calculated by dividing the atmospheric dispersion values (Chi/Q) for 6652 by the atmospheric dispersion values for the 300 Area. The code CAP88-PC was used to calculate these dispersion values. Similar calculations were performed to obtain location correction factors for other Pacific Northwest-operated DOE facilities outside of the 300 Area.

2.3 Potential Emission Dose Assessment

Doses from projected radionuclide emissions were calculated by multiplying the quantity of each radionuclide present in the facility by its associated potential release fraction, the 300 Area unit dose factor (see Appendix C), and the location correction factor. Doses from individual radionuclides were summed to derive the total potential annual emission dose for each facility.

The assessments are independently reviewed, then packets are prepared for each building containing the raw inventory information, any communications clarifying or correcting the inventory information, summarized inventory information, and a cover sheet showing the resulting dose and approval signatures. The packets are approved by the preparer, the reviewer, the divisional points of contact for the inventory custodians, and the building managers. Once approved, assessment packets are maintained as records by the Effluent Management Group. A summary of the results of the assessment for 1999 and the primary contributing radionuclides for facilities that require continuous monitoring is provided in Table 4. The table also identifies the radionuclides that contribute 10% or more of the potential dose for facilities that are required to be continuously sampled.

Table 3. Unit Dose Factors for 300 Area Source Location

Nuclide	Dose (mrem/Ci)		Nuclide	Dose (mrem/Ci)	Nuclide	Dose (mrem/Ci)
Ac-225	3.10E-00		Co-57	6.10E-02	Kr-89	7.60E-05
Ac-227	3.30E+02	l	Co-58	1.20E-01	Kr-90	3.80E-08
Ac-228	3.80E-02		Co-60	5.00E-00	La-140	9.30E-03
Ag-109m	5.50E-10	\vdash	Cr-51	2.30E-03	La-141	1.80E-05
Ag-110	6.40E-11	\vdash	Cs-134	2.10E-00	La-142	9.20E-04
Ag-110m	9.60E-01	<u> </u>	Cs-134m	2.90E-05	Mn-54	3.40E-01
Ag-111	1.90E-02		Cs-135	1.20E-01	Mn-56	8.70E-04
Am-241	2.90E+02		Cs-136	8.80E-02	Mo-93	5.60E-02
Am-242	2.70E-02	-	Cs-137	4.90E-00	Mo-99	5.20E-03
Am-242m	2.80E+02		Cs-138	5.90E-04	N-13	1.30E-04
Am-243	2.90E+02		Cs-139	4.00E-05	Na-22	2.40E-00
Ar-41	3.30E-04		Cu-64	3.40E-04	Na-24	4.50E-03
As-76	3.00E-03		Eu-152	4.90E-00	Nb-93m	2.80E-02
Ba-133	1.50E-00		Eu-152m	2.30E-04	Nb-94	1.60E+01
Ba-133m	8.60E-04		Eu-154	4.00E-00	Nb-95	6.40E-02
Ba-133m	1.50E-05		Eu-155	1.70E-01	Nb-95m	4.60E-03
Ba-137III Ba-139	8.80E-05		Eu-156	7.40E-02	Nb-97	2.50E-04
Ba-139			F-18	3.60E-04	Nb-97m	
	7.90E-02		Fe-55	6.90E-03	Nd-147	9.40E-07
Ba-141 Ba-142	1.60E-04				 	2.20E-02
	1.20E-04		Fe-59	1.20E-01	Ni-59	8.10E-03
Be-7	4.50E-03		Fr-221	4.90E-04	Ni-63	1.00E-02
Bi-210 Bi-211	9.90E-02 3.20E-05		Fr-223	8.30E-04	Ni-65	3.50E-04
			Ga-67	1.70E-03	Np-237	2.60E+02
Bi-212	1.30E-02		H-3	4.10E-04	Np-238	1.70E-02
Bi-213	4.90E-04		Hf-181	7.00E-02	Np-239	3.40E-03
Bi-214	4.10E-04 6.30E-03		Hg-203 Ho-166	4.40E-02 2.10E-03	Np-240	3.70E-04 3.70E-05
Br-82	2.70E-06		Ho-166m	1.60E+01	Np-240m O-15	1.50E-05
Br-83 Br-84	4.30E-04		I-122	2.50E-05	P-32	5.70E-02
	2.10E-06		I-123	1.20E-03	Pa-231	
Br-85 C-11	2.10E-06 1.90E-04		I-125	2.80E-00	Pa-231 Pa-233	2.70E+02 3.50E-02
C-11 C-14			<u> </u>	4.80E+01	Pa-234	
	5.00E-02		I-129			1.70E-03
Ca-41	2.00E-04 1.20E-01	*	I-130	1.30E-02 1.40E-00	Pa-234m Pb-209	4.50E-08
Cd-109		ļ.,	I-131			4.30E-05
Cd-115	5.20E-03			2.50E-03	Pb-210	6.30E+01
Cd-115m	1.40E-01		I-133	1.00E-02	Pb-211	3.40E-03
Ce-141	2.70E-02	-	I-134	1.10E-03	Pb-212	8.10E-02
Ce-143	2.80E-03		I-135	4.60E-03	Pb-214	3.60E-04
Ce-144	3.70E-01		In-113m	1.00E-04	Pd-107	8.50E-03
Cf-252	7.90E+01	*	In-115 In-115m	2.00E-00	Pd-109	7.60E-04
Cl-36	2.20E-01	Ϊ	 	1.50E-04	Pm-147	2.80E-02
Cm-242	9.10E-00		Ir-192	1.40E-01	Pm-148	3.00E-02
Cm-243	1.90E+02	<u> </u>	K-40	3.30E-00	Pm-148m	2.00E-01
Cm-244	1.50E+02		Kr-83m	6.00E-08	Pm-149	3.20E-03
Cm-245	3.00E+02		Kr-85	9.20E-07	Pm-151	6.00E-04
Cm-246	2.90E+02	 	Kr-85m	4.30E-05	Po-210	2.10E+01
Cm-247	2.70E+02	<u> </u>	Kr-87	2.10E-04	Po-218	3.10E-06
Cm-248	1.10E+03		Kr-88	5.90E-04	Pr-143	2.50E-02

Nuclide	Dose (mrem/Ci)	Nuclide	Dose (mrem/Ci)	Nuclide	Dose (mrem/Ci)
Pr-144	2.80E-05	Sn-126	9.00E-01	Xe-135	6.70E-05
Pr-144m	6.30E-06	Sr-89	7.60E-02	Xe-135m	6.80E-05
Pu-236	4.40E+01	Sr-90	4.90E-00	Xe-137	9.80E-06
Pu-238	1.80E+02	Sr-91	1.00E-03	Xe-138	1.90E-04
Pu-239	2.00E+02	Sr-92	7.80E-04	Y-90	1.10E-02
Pu-240	2.00E+02	Tb-160	1.60E-01	Y-90m	2.60E-04
Pu-241	3.20E+02	Tc-101	5.20E-05	Y-91	9.80E-02
Pu-242	1.80E+02	Tc-97	1.30E-01	Y-91m	1.50E-04
Pu-243	1.10E-04	Tc-99	6.20E-01	Y-92	6.30E-04
Pu-244	1.80E+02	Te-125m	3.50E-02	Y-93	1.30E-03
Ra-223	6.60E-00	Te-127	1.90E-04	Zn-65	4.30E-01
Ra-224	2.00E-00	Te-127m	8.50E-02	Zn-69	2.60E-05
Ra-225	4.70E-00	Te-129	6.80E-05	Zn-69m	8.80E-04
Ra-226	1.70E+01	Te-129m	8.60E-02	Zr-93	3.20E-02
Ra-228	7.90E-00	Te-131	1.20E-04	Zr-95	1.30E-01
Rb-86	7,20E-02	Te-131m	5.20E-03		
Rb-87	2.60E-01	Te-132	1.30E-02		-
Rb-88	1.50E-04	Te-133	1.40E-04		
Rb-89	3.70E-04	Te-133m	6.00E-04		
Rb-90	6.60E-05	Te-134	2.10E-04		
Rb-90m	2.10E-04	Th-227	5.50E-00		
Re-187	6.40E-04	Th-228	1.20E+02		
Rh-103m	4.90E-00	Th-229	3.20E+02		
Rh-105	9.40E-04	Th-230	1.20E+02		
Rh-105m	7.80E-09	Th-231	5.40E-04		
Rh-106	2.80E-09	Th-232	1.90E+02		
Rn-219	1.40E-34	Th-234	9.90E-02		7
Rn-220	5.50E-07	Tl-207	1.10E-06		
Rn-222	1.20E-03	T1-208	1.40E-04		
Ru-103	5.00E-02	TI-209	3.90E-05		
Ru-105	6.60E-04	U-232	2.40E+02		
Ru-106	4.70E-01	U-233	7.10E+01		
Ru-97	1.60E-03	U-234	7.00E+01		
S-35	6.20E-03	U-235	6.70E+01		
Sb-124	2.20E-01	U-236	6.60E+01		
Sb-125	5.40E-01	U-237	1.10E-02		
Sb-126	9.10E-02	U-238	6.20E+01		
Sb-126m	3.00E-04	U-240	1.10E-03		
Sb-127	1.40E-02	W-181	9.50E-03		
Sc-46	2.70E-01	W-185	1.50E-02		
Se-75	9.10E-01	W-187	1.10E-03		
Se-79	1.70E-00	* Xe-122	1.80E-05		
Sm-147	3.60E+01	Xe-123	1.60E-04		
Sm-151	1.80E-02	Xe-125	6.80E-05		
Sm-153	2.10E-03	Xe-127	7.20E-05		
Sn-113	4.00E-02	Xe-131m	2.60E-06		
Sn-123	1.00E-03	Xe-133	9.60E-06		
Sn-125	5.70E-02	Xe-133m	8.40E-06		

^{*} Calculated using the GENII Code.

 Table 4. Emission System Potential Dose Assessment Summary

Emission	Emission	System Description	Emission	Nuclides	Comment
System	Type ¹		Measurement	Contributing >	
			Required	10% of Potential	
			-	Dose	
	S	ystems Located in the Southeas	t Region (300 Area)) of the Hanford Site	
300-N	Fugitive	Caisson	None	None	Sealed
,					Sources Only
305-B	Fugitive	Hazardous Waste Storage	Confirmatory	Pu-238	
	+ Point	Facility			
306-W	Point	Materials Development	Confirmatory	U(20%),	
		Building		U(depleted)	
314	Fugitive	Engineering Development	None	U(natural)	Bldg vacated
	+ Point	Laboratory			(Inventory
					based on
					holdup)
318	Point	Radiological Calibrations	None	Eu-154, U(natural)	Primarily
		Laboratory			sealed sources
320	Point	Analytical and Nuclear	Confirmatory	U-233	
	1	Research Laboratory			
323	Point	Mechanical Properties	Confirmatory	Co-60, V-49	
		Laboratory	•	,	
325	Point	Radiochemical Processing	Continuous	H-3, Pu-238	
		Laboratory		11-5, 1 u-250	
326	Point	Materials Sciences	Confirmatory	Gd-149, U(natural)	
		Laboratory			
329	Point	Chemical Sciences	Confirmatory	Am-241	
		Laboratory	,		·
331	Point	Life Sciences Laboratory I	Continuous	Pu-339, Pu-241	
331G	Point	Interim Tissue Repository	Confirmatory	C-14	
331H	Point	Aerosol Wind Tunnel	None	None	Sealed
		Research Facility			Sources Only
3718A	Fugitive	Lab Equipment Central Pool	None	U(natural)	
3720	Point	Environmental Sciences	Continuous	Am-243, Pu-238	1999 PTE <
		Laboratory			0.1 mrem;
			1		facility in
					transition
3730	Point	Gamma Irradiation Facility	Confirmatory	Co-60, V-49	
3745	Point	Radiological Calibrations	Confirmatory	U(Hanford)	
		and Standards Building			
3020	Point	Environmental Molecular	None	None	Sealed
EMSL		Sciences Laboratory			Sources Only

200E	Fugitive	Environmental Monitoring Shed	None	None	Sealed Sources Only
FLTF	Fugitive	Field Lysimeter Test Facility	None	None	Sealed Sources Only
PSB	Fugitive	Prototype Surface Barrier	None	None	Sealed Sources Only
2718E	Fugitive	Critical Mass Fissile Storage	None	None	Radioactive Material in DOT containers
622R	Fugitive	Meteorological Laboratory	None	None	Sealed Sources Only
		Systems 1	Located on ALE		
6652	Point	Arid Land Ecology Laboratory	None	None	Sealed Sources Only
		Systems Lo	cated in Richlan	đ	
74 7 A	Fugitive	Whole Body Counter	None	None	Primarily Sealed Sources

¹ "Fugitive emissions" are radioactive air emissions that do not and could not reasonably pass through a stack, vent, or other functionally equivalent structure, and that are not feasible to directly measure and quantify (WAC 246-247).

3.0 References

10 CFR 835 1998. "Occupational Radiation Protection." U.S. Environmental Protection Agency, Washington, D.C.

40 CFR 61 1989. "National Emission Standards for Hazardous Air Pollutants." U.S. Environmental Protection Agency, Washington, D.C.

Ballinger, M. Y., S. J. Jette, and M. J. Sula. 1995. Assessment of Unabated Facility Emission Potentials for Evaluating Airborne Radionuclide Monitoring Requirements at Pacific Northwest National Laboratory - 1995. PNL-10855, Pacific Northwest Laboratory, Richland, Washington.

DOE. 1988. General Environmental Protection Program, DOE 5400.1, U.S. Department of Energy Order. U.S. Department of Energy, Washington, D.C.

DOE. 1991. Environmental Regulatory Guide for Radiological Effluent Monitoring and Environmental Surveillance, DOE/EH-0173T. U.S. Department of Energy, Washington, D.C.

EPA. 1990. National Emission Standards for Hazardous Air Pollutants. 40 CFR 61. Environmental Protection Agency, U.S. Code of Federal Regulations.

Napier, B. A., R. A. Peloquin, D. L. Strenge, and J. V. Ramsdell. 1988. GENII - The Hanford Environmental Radiation Dosimetry Software System. PNL-6584, Vols. 1-3. Pacific Northwest Laboratory, Richland, Washington.

Pacific Northwest National Laboratory. 1999a. Facility Effluent Monitoring Plan for the 325 Radiochemical Processing Laboratory. PNNL-12157. Pacific Northwest National Laboratory, Richland, Washington.

Pacific Northwest National Laboratory. 1999b. Facility Effluent Monitoring Plan for the 331 Complex. PNNL-12158. Pacific Northwest National Laboratory, Richland, Washington.

Pacific Northwest National Laboratory. 1999c. Facility Effluent Monitoring Plan for the 3720 Building. PNNL-12159. Pacific Northwest National Laboratory, Richland, Washington.

Pacific Northwest National Laboratory. 1999d. Facility Effluent Monitoring Plan for the Balance-of-Plant Facilities. PNNL-12160. Pacific Northwest National Laboratory, Richland, Washington.

Parks, B. S. 1992. *User's Guide for CAP88-PC Version 1.0.* 402-B-92-001, U.S. Environmental Protection Agency, Office of Radiation Programs, Las Vegas Facility, Las Vegas, Nevada.

Sula, M. J. and S. J. Jette. 1994. Pacific Northwest Laboratory Facilities Radionuclide

Inventory Assessment CY 1992-1993. PNL-10061. Pacific Northwest Laboratory, Richland, Washington.

WAC. 1994. *Radiation Protection - Air Emissions*, WAC 246-247, Washington Administrative Code. Washington Department of Health.

Westinghouse Hanford Company. 1991. *Unit Dose Calculation Methods and Summary of Facility Effluent Monitoring Plan Determinations*. WHC-EP-0498. Westinghouse Hanford Company, Richland, Washington.

Appendix A

Common Radionuclide Mixtures

Weapons Grade (6% Pu-240)	Isotope	wt fraction	Isotope	Mixture
Weapons Grade (0701 a 240)	Isotope	we made on	Ci/g	Ci/g
	Pu-238	0.0005	17.5	0.01
	Pu-239	0.93	0.062	0.01
	Pu-240	0.061	0.002	0.00
	Pu-242	0.0005	0.0039	0.00
	Am-241	0.0003	3.24	0.00
	AIII-241	<u> </u>	3.24	8.02E-02
		· .		8.02E-02
Fuel Grade (12% Pu-240)	Isotope	wt fraction	Isotope	Mixture
1 uel Glade (12/81 u-240)	Isotope	willaction	Ci/g	Ci/g
•	Pu-238	0.001	17.1	0.02
	Pu-238 Pu-239	0.001	0.062	0.02
	Pu-240	0.124	0.226	0.03
	Pu-242	0.001	0.0039	0.00
	Am-241	0	3.24	0.00
				9.75E-02
24% Pu-240	Isotope	wt fraction	Isotope	Mixture
			Ci/g	Ci/g
	Pu-238	0.018	17.1	0.31
	Pu-239	0.542	0.062	0.03
•	Pu-240	0.238	0.226	0.05
·	Pu-242	0.064	0.0039	0.00
	Am-241	0	3.24	0.00
				3.95E-01
Depleted U	Isotope	wt. Fraction	Isotope	Mixture
			Ci/g	Ci/g
	U-234	4.2E-06	6.20E-03	2.60E-08
	U-235	0.0045	2.10E-06	9.45E-09
	U-238	9.95E-01	3.30E-07	3.29E-07
				3.64E-07

Natural U	Isotope	wt. Fraction	Isotope	Mixture
			Ci/g	Ci/g
	U-234	5.5E-05	6.20E-03	3.41E-07
	U-235	0.0072	2.10E-06	1.51E-08
	U-238	9.93E-01	3.30E-07	3.28E-07
				6.84E-07
		·		
Enriched U (<20%)	Isotope	wt. Fraction	Isotope	Mixture
			Ci/g	Ci/g
i	U-234	0.0014	6.20E-03	8.68E-06
	U-235	0.2	2.10E-06	4.20E-07
	U-238	0.7986	3.30E-07	2.64E-07
			3	9.36E-06
			·	
Enriched U (<90%)	Isotope	wt. Fraction	Isotope	Mixture
			Ci/g	Ci/g
	U-234	0.0097	6.20E-03	6.01E-05
	U-235	0.9	2.10E-06	1.89E-06
	U-238	0.0903	3.30E-07	2.98E-08
				6.21E-05

Reference for Pu and U mixes isotope weight percent: Sula, M.J., E.H. Carbaugh, and D.E. Bihl. 1991. Technical Basis for Internal Dosimetry at Hanford, PNL-6866, Rev. 1, Pacific Northwest Laboratory, Richland, Washington.

Appendix B

Radionuclide Inventory Database Features

The information obtained from the nuclear materials inventory, composite radioactive materials inventory, and facility management radioactive materials inventory is maintained by Effluent Management (EM) in an ACCESS database. The inventory information that is obtained and stored in the database is listed in Section 2.1.

Database Population

The nuclear materials and composite radioactive materials inventories currently are obtained in electronic formats. The general format of the electronic files is manipulated to allow electronic downloading of the information from these two sources into the database tables, the data in the electronic files is not changed.

The facility management radioactive materials inventory is obtained from individual research personnel who act as custodians of the material. This inventory data is provided in various forms (handwritten, electronic files, e-mail messages, etc.) Electronic files are reformatted to the database table format and downloaded directly into the database tables. Handwritten and e-mail information is manually input to the database tables by EM personnel. Entries to the database are independently verified by other EM personnel for accuracy.

The information entered into the database includes:

- Staff member acting as custodian of the material
- Research or support division of the custodian
- Form of the material gas, liquid, powder, solid, contained, or exempt/sealed DOT
- Basis for the Inventory physical inventory records, estimated through process knowledge, estimated through transportation records, or estimated through other means (i.e., procedures, documents, etc.)
- Nuclide
- Inventory in activity (Ci) or mass (grams)
- Building/Room in which material is stored or used
- Identification number numbers previously assigned by the custodian or by other databases (sealed source or material balance area identification numbers)
- Comments any additional comments related to the material (e.g., reference numbers on the material, whether or not the material is considered throughput, description of the material, etc.).

The database assigns an identification number to each of the entries. For sealed source and Material Balance Area (MBA) data the identification number previously assigned to the material is used. For the research inventories, the database applies a numerical sequential number to each

of the entries. The database assigns the following designations to the identification number:

D - = Input provided by the research division

G -= Government sealed source

P - = Private radioactive material inventory

M - = Nuclear materials inventory.

Other Database Tables

Other database tables are used to store specific information about individual nuclides that are used in the NESHAP dose calculations:

Factors Table - Factors for the dose per curie (mrem/Ci) for different isotopes are listed in Table 3 and are entered as a separate table in the ACCESS database. The source for each of the nuclide factors is listed in the factors table.

Release Fraction Table - The release fractions for material forms (i.e. solid, liquid, powder, etc.) are listed in Table 2. These are entered as a separate table in the database.

Calculations

The ACCESS database uses queries and macros that are applied to the tables listed above to calculate the potential dose for the different Pacific Northwest facilities.

Normalizing Inventory Data -- The database is designed to convert the reported mass and activity inventory units (i.e., grams, mCi, μ Ci, mg, μ g, etc.) to curie (Ci) units for use in subsequent calculations.

Potential Dose Calculations -- Potential dose calculations are determined on a facility-specific basis. The reported inventory is first converted to Ci and then it is multiplied by the dose factor (mrem/Ci) for the specific nuclide, the location modification factor, and the release fraction to determine the potential dose for that nuclide inventory.

Example:

20 µCi of U-238 in powder form at the 305B facility

 $20 \,\mu\text{Ci} * 1.0\text{E} - 06 \,\text{Ci}/\mu\text{Ci} * 1\text{E} - 03 * 6.2\text{E} + 01 \,\text{mrem/Ci} * 1.0 = 1.24\text{E} - 06 \,\text{mrem/Ci}$

Where:

1E-03 = release fraction for powder

6.2E+01 mrem/Ci = dose factor for U-238

1.0 = location modification factor (with relation to 300 Area)

The cumulative dose for the facility is determined by summing the potential doses of each inventory entry.

Reports

Three reports are generated using the database for the NESHAP assessment as part of the final packet:

- The first is a report of the "raw" data that was provided by the research personnel, the nuclear materials inventory, and the composite radioactive materials inventory.
- The second is a summary page listing the potential dose (mrem/Ci) of the facility inventory and sign-off blocks for the preparer, reviewer, facility safety representative, divisional representative(s), and the building manager.
- The third is a listing of each radionuclide present in the building and the dose contribution in mrem/Ci and in % of the total dose.

Appendix C

Unit Dose Factor Calculations



Project No.	
-------------	--

Internal Distribution

MJ Sula

File/LB

K Rhoads

Date

June 3, 1996

To

MY Ballinger

From

LH Staven

Subject

331 Building Unit Ci Release Dose Calculations

Please find attached the Unit Ci dose calculations for the 331 Building. These calculations have been reviewed by Kathy Rhoads for HEDOP approval. The 331 values are reported with the 9 year meteorological data.

These values were calculated based on the EMSL values reported earlier, and are adjusted by 331 specific X/Q values. During the preparation of these calculations, USGS quadrangle maps were made available for determining distances to the maximally exposed individuals. These maps will assure consistency in reporting dose conversion values. Please use these values for your reporting purposes for the 331 Building.

If you have any questions, please contact me at (509) 375-2429.

Attachment

HEDOP REVIEW CHECKLIST for Radiological and Nonradiological Release Calculations

Document reviewed (include title or description of calculation, document number, author, and date, as applicable):

CAP88-PC Dose Conversion Factors for 331 Building 5/22/96

Submitted by:	L. H. Staven	Date Submitted:	5/22/96
Scope of Review:			
YES NO* N/A			
[x] []	environmental transp	review and approval opert and dose calculati	on portion of
[x] [] []	2. Detailed technical n	en performed and docume review(s) and approval(nations have been perfo	s) of scenario
[x] [] [] []	3. HEDOP-approved code	(s) were used. were selected according	to HEDOP
[x]. [] []	5. All applicable envir	ronmental pathways and re appropriate for the	
[x] [] [x]	6. Hanford site data we	ere used. Kternal to the computer	
[x] [] [x]	 The analysis is cons Supporting notes, caresolutions, or other 		comment hed. (Use the
[x] []	10. Approval is granted Environmental Dose (rd
* All "NO" respo	nses must be explained ar	nd use of nonstandard m	ethods
<u>K. Rhoads</u> HEDOP-Approv	Khowh red Reviewer (Printed Name	e and Signature)	5/30/96 Date
	itional signed and dated		

APPLICATION REPORT

- 1) Project title and number <u>CAP88-PC DOSE CONVERSION FACTORS FOR 331</u> <u>BUILDING 5/22/96</u>
- 2) Purpose of application package and relationship to other work:

This series of CAP88-PC runs were produced in order to report the 50-year total effective dose equivalent (TEDE) to the maximally exposed individual (MEI) and offsite worker per Ci released from the 331 Building, Table 1. These values will be used to demonstrate NESHAPs (National Emissions Standards for Hazardous Air Pollutants) compliance.

This application is based on the work performed in January '96 for the EMSL facility. The dose factors for the EMSL facility were multiplied by the appropriate conversion factor. Each Conversion factor is specific to the location of the MEI of interest, and is determined by the ratio of the (331 MEI/EMSL MEI) chi/Q values.

3) List original sources of input data, assumptions and derivations used to obtain it, and justification for its use, as appropriate. (If input information has been previously reviewed, reference the documentation of this review.)

Dose factors were created for both the offsite MEI and the onsite MEI.

Worker doses were calculated based on 100% import of food products. To adjust for 2000 hours per year at the site, the CAP88 PC dose was adjusted by 2000/8766.

331 Building parameters included a 10 m release height.

Hanford parameters listed in Schreckhise et al (1993) were used in place of default CAP88-PC default parameters to better model the Hanford environment.

- 4) Minor changes made in the software that produced the application run (see section 4.1). N/A
- 5) Describe interrelationships and dependencies of each application run in the application package.

The TEDEs for the CAP88-PC radionuclides were adjusted to represent the value with the contribution from progeny which were grown-in after the release from the facility. The adjustment factors were derived from previous GENII runs (FEMP 3/93) where only the parent is released, and the output includes the doses from the progeny. The adjustment factor was taken by dividing the dose from the parent by the total dose from the parent and progeny. The CAP88-PC value was then divided by the factor to represent the total dose. Where CAP88-PC

calculated chains (i.e., Cs-137) the progeny were added to the parent, and no other adjustment was made.

6) Summarize the overall output of the application package in relation of the purpose stated in item 2 above (including tables and graphs, as appropriate):

Table 1 shows the worker and resident 50 year TEDE conversion factors per Ci released.

Date

7)	Submitted for ITR by:	
	Sol VII	, ,
	X desta-	5/22/96
	LH Staven	Date

8) Approved for reporting results to sponsor by:

APPLICATION RECORD LOG

1) Project title and number: <u>CAP88-PC Dose Conversion Factors for EMSL Building</u>

2)Application 3) Date and Time 4)Software Name (title) 5) computer type Record Log of Run and Version

6) Comments Identifier operating system

7)Data input

Number

compilers/libraries

Filename (a) see below

CAP88-PC Ver 1.0

Gateway2000 486

see below

see below

Directory of D:\CAP88PC\OUTPUT

Directory of D:\CAP88PC\OUTPUT

Worker

05/22/96 14:33 .SUM 331rn wrkr onsit 05/22/96 14:27 331rn wrkr offsi .SUM

Resident

9635 05/07/96 12:21 .SUM 331rn resid

Prepared by: LASta 5/22/96

CAP88PC	ose Factor	s for Unit C	Releases	From 331 F	uilding	1 1	
1	1	1	1	1	1	1 - 1	-
	 		<u> </u>	<u> </u>	ļ	 	
	fraction of	Officia	<u> </u>	0-24-	· · · · · · · · · · · · · · · · · · ·	W	
				Onsite		5	Ļ_
	dose from		ļ	Worker		Residentia	
X/Q =	parent	2.82E-06		1.49E-05		1.77E-06	
Location	ļ	1000 m N\	<u> </u>	200 m S		1000 m E	
Duration		2000 h/y		2000 h/y		8766 h/y	
d locally pro	duced	0%		0%		100%	
		mrem/yr		mrem/yr		mrem/yr	
Radionucl	ide	per Ci rele	ased	per Ci rele	ased	per Ci rele	ase
		F					
H-3		2.3E-05		1.22E-04		4.1E-04	_
BE-7		1.3E-03		6.87E-03		4.5E-03	├-
C-11		7.2E-05		3.80E-04		1.9E-04	-
C-14	ļ						-
		2.1E-06		1.12E-05		5.0E-02	<u> </u>
N-13		4.8E-05		2.55E-04		1.3E-04	_
0-15		5.6E-06		2.96E-05		1.5E-05	L_
F-18		1.3E-04		7.08E-04		3.6E-04	
NA-22		8.4E-01		4.45E+00		2.4E+00	L
NA-24		1.7E-03		8.80E-03		4.5E-03	
P-32		1.0E-03		5.31E-03		5.7E-02	
S-35		4.9E-05		2.61E-04		6.2E-03	_
AR-41	 	1.2E-04		6.49E-04		3.3E-04	-
K-40		5.0E-01		2.64E+00		3.3E+00	
CA-41	ļ	7.5E-05		3.97E-04		2.0E-04	<u> </u>
SC-46		7.8E-02		4.14E-01		2.7E-01	
CR-51		4.8E-04		2.55E-03		2.3E-03	
MN-54		1.2E-01		6.21E-01		3.4E-01	
MN-56		3.2E-04		1.72E-03		8.7E-04	
FE-55		3.2E-04		1.67E-03		6.9E-03	
FE-59		2.4E-02		1.29E-01		1.2E-01	_
CO-57		1.8E-02		9.70E-02		6.1E-02	
CO-58	l	3.3E-02		1.77E-01		1.2E-01	
CO-60		1.8E+00		9.32E+00		5.0E+00	_
							_
NI-59		1.7E-03		8.84E-03		8.1E-03	
NI-63		3.8E-04		2.01E-03		1.0E-02	
NI-65		1.3E-04		6.87E-04		3.5E-04	_
CU-64		1.2E-04		6.52E-04		3.4E-04	
ZN-65		6.3E-02		3.34E-01		4.3E-01	
ZN-69		9.6E-06		5.07E-05		2.6E-05	
ZN-69M		3.2E-04		1.70E-03		8.8E-04	
GA-67		3.5E-04		1.86E-03		1.7E-03	
AS-76		1.0E-03		5.28E-03		3.0E-03	
BR-82		2.2E-03		1.16E-02		6.3E-03	
	 			5.28E-06		2.7E-06	
BR-83		1.0E-06					<u> </u>
BR-84	<u> </u>	1.6E-04		8.49E-04	<u> </u>	4.3E-04	_
BR-85		7.9E-07		4.18E-06	· · · · · · · · · · · · · · · · · · ·	2.1E-06	ļ
KR-83M	•	2.2E-08		1.17E-07	*	6.0E-08	
KR-85		3.4E-07		1.81E-06		9.2E-07	
KR-85M		1.6E-05		8.35E-05		4.3E-05	
KR-87		8.0E-05		4.21E-04		2.1E-04	
KR-88		2.2E-04		1.15E-03	<u> </u>	5.9E-04	
KR-89		2.8E-05		1.50E-04		7.6E-05	
KR-90	<u> </u>	1.3E-08		6.94E-08		3.5E-08	_
RB-86		2.1E-03		1.13E-02		7.2E-02	<u> </u>
RB-87		1.2E-03		6.49E-03		2.6E-01	<u></u>
RB-88		5.7E-05		3.01E-04		1.5E-04	<u></u>
RB-89		1.4E-04		7.35E-04		3.7E-04	L
RB-90		2.4E-05		1.29E-04		6.6E-05	
RB-90M		7.8E-05		4.14E-04		2.1E-04	

^{*} Includes dose from long-lived progeny
** Includes dose from short-lived progeny

CAPSSPC	Dose Factor	s for Unit Ci F	Releases From 331 B	uilding	
<u> -</u>	fraction of		Onsite		
	dose from		Worker		lential
X/Q =	parent	2.82E-06	1.49E-05		E-06
Location		1000 m NW	200 m S	1000	
Duration		2000 h/y	2000 h/y	8766	
od locally pr	oduced	0%	0%	•	100%
		mrem/yr	mrem/yr	mren	
Radionuc	lide	per Ci releas	ed per Ci rele	ased per C	i released
SR-89		1.1E-03	5.73E-03	7.6	E-02
SR-90 *	0.94	3.9E-02	2.07E-01		E+00
SR-91		3.7E-04	1.94E-03	1.0	E-03
SR-92		2.9E-04	1.53E-03	7.8	E-04
Y-90		1.6E-03	8.25E-03	1.1	E-02
Y-90M		9.7E-05	5.11E-04	2.6	E-04
Y-91		8.2E-03	4.35E-02	9.8	E-02
Y-91M		5.6E-05	2.94E-04		E-04
Y-92		2.3E-04	1.23E-03	6.3	E-04
Y-93		4.9E-04	2.58E-03	1.3	E-03
ZR-93	1	8.0E-03	4.21E-02		E-02
ZR-95 *	0.75	3.2E-02	1.69E-01		E-01
NB-93M		6.5E-03	3.45E-02		E-02
NB-94		5.8E+00	3.09E+01		E+01
NB-95	1	1.3E-02	7.08E-02		E-02
NB-95M		5.3E-04	2.80E-03		E-03
NB-97		9.2E-05	4.87E-04		E-04
NB-97M		3.5E-07	1.84E-06		E-07
MO-93		2.1E-02	1.10E-01		E-02
MO-99 **		1.2E-03	6.12E-03		E-03
TC-99M	-	0.0E+00	0.00E+00		E+00
TC-97	+	2.3E-02	1.23E-01		E-01
TC-99	 	1.4E-03	7.39E-03		E-01
TC-101		1.9E-05	1.03E-04		E-05
RU-97	-	4.3E-04	2.26E-03		E-03
RU-103		1.0E-02	5.52E-02		E-03
RU-105	-	2.4E-04	1.29E-03		E-04
RU-106		7.9E-02	4.18E-01		E-01
RH-105	-	2.4E-04	1.29E-03		E-04
RH-105M	-	2.4E-04 2.9E-09	1.54E-08		E-04
RH-106	-	1.0E-09	5.45E-09		E-09
PD-107		2.1E-03	1.11E-02		E-03
PD-107		2.1E-03 2.7E-04	1.43E-03		E-04
			1.43E-03 1.09E-09		E-10
AG-109M	<u> </u>	2.1E-10 2.4E-11			
AG-110			1.25E-10		E-11
AG-110M	' 	3.1E-01	1.65E+00		E-01
AG-111		1.2E-03	6.28E-03		E-02
CD-115		1.0E-03	5.31E-03		E-03
CD-115M	<u> </u>	7.7E-03	4.07E-02		E-01
IN-113M		3.8E-05	2.03E-04		E-04
IN-115		1.7E-01	9.01E-01		E+00
IN-115M		5.7E-05	3.02E-04		E-04
SN-113		2.6E-03	1.39E-02		E-02
SN-123		3.8E-04	2.01E-03		E-03
SN-125		4.0E-03	2.11E-02		E-02
SN-126		2.3E-01	1.23E+00		E-01
SB-124		5.2E-02	2.77E-01		E-01
SB-125		1.9E-01	1.00E+00		E-01
SB-126		1.8E-02	9.56E-02	0.1	E-02

^{*} Includes dose from long-lived progeny
** Includes dose from short-lived progeny

		f	0%-:4-		0				
		fraction of dose from		ļ ———	Onsite Worker	 	Residentia	<u> </u>	
	X/Q =				1.49E-05				
		parent	2.82E-06 1000 m NV		200 m S		1.77E-06 1000 m E		
	Location	 	2000 h/y	1				ļ	ł
Foo	d locally pro	ducod	2000 h/y	ļ	2000 h/y		8766 h/y 100%		1
F00	d locally pro	Tucea	mrem/yr		mrem/yr				
	Radionucl	ide	per Ci rele	acad	per Ci rele		mrem/yr per Ci rele	2001	
	Tadionaci	100	per Orrese	aseu	per Orreie	aseu	per Oriele	aseu	-
	SB-126M		1.1E-04		5.83E-04		3.0E-04		
	SB-127		2.3E-03		1.23E-02		1.4E-02		
	TE-125M		1.8E-03		9.39E-03		3.5E-02		
	TE-127		7.1E-05		3.76E-04		1.9E-04		
	TE-127M		3.9E-03	1	2.07E-02		8.5E-02		, .
	TE-129		2.5E-05		1.34E-04		6.8E-05		
	TE-129M		4.6E-03		2.41E-02		8.6E-02		İ
	TE-131		4.6E-05		2.42E-04		1.2E-04		
	TE-131M		1.7E-03		9.11E-03		5.2E-03	1	
	TE-132	1	1.7E-03		9.11E-03		1.3E-02		
	TE-133	I	5.0E-05		2.66E-04		1.4E-04		l
	TE-133M		2.2E-04		1.18E-03		6.0E-04		.
	TE-134		7.8E-05		4.11E-04		2.1E-04		į
	I-122		9.3E-06		4.90E-05		2.5E-05		
	I-123		4.1E-04		2.18E-03		1.2E-03		
	1-125		7.6E-03		4.00E-02		2.8E+00		
	I-129		6.9E-01		3.66E+00		4.8E+01		
	1-130		4.6E-03		2.45E-02		1.3E-02		
	I-131		1.5E-02		8.08E-02		1.4E+00		
	I-132		9.3E-04		4.90E-03		2.5E-03		
	I-133		2.6E-03		1.35E-02		1.0E-02		
	I-134		4.2E-04		2.23E-03		1.1E-03	1	
	I-135		1.7E-03		9.01E-03		4.6E-03		
	XE-122		6.9E-06		3.62E-05	1	1.8E-05		İ
	XE-123		6.0E-05		3.15E-04		1.6E-04		
	XE-125		2.5E-05		1.33E-04		6.8E-05	1	
	XE-127		2.7E-05		1.40E-04		7.2E-05		
	XE-131M		9.7E-07		5.14E-06		2.6E-06		
	XE-133		3.6E-06		1.88E-05		9.6E-06		
	XE-133M		3.1E-06		1.65E-05		8.4E-06		
<u> </u>	XE-135	<u> </u>	2.5E-05		1.31E-04		6.7E-05		
	XE-135M		2.5E-05		1.33E-04		6.8E-05		İ
<u> </u>	XE-137		3.6E-06		1.92E-05	1	9.8E-06		
	XE-138 CS-134	 	7.1E-05 5.2E-01		3.73E-04 2.73E+00		1.9E-04 2.1E+00		
	CS-134 CS-134M	 	1.1E-05		5.63E-05		2.1E+00 2.9E-05		
	CS-134M CS-135	 	7.8E-04		4.11E-03		1.2E-01		
	CS-135	 	1.4E-02		7.52E-02		8.8E-02		
	CS-136		1.4E-02 1.6E+00		8.34E+00		4.9E+00		_ Bau
	- BA-137M		0.0E+00		0.00E+00		0.0E+00	<u> </u>	- 1300 W
	CS-138		2.2E-04		1.15E-03		5.9E-04	 	
	CS-139	 	1.5E-05		7.77E-05		4.0E-05		1
1	BA-133	 	5.5E-01		2.90E+00		1.5E+00		
+-	BA-133M		1.6E-04		8.63E-04		8.6E-04		
	BA-137M	 	5.5E-06		2.93E-05		1.5E-05		
	BA-139		3.3E-05		1.73E-04		8.8E-05	1	
	BA-140 **	 	1.4E-02		7.54E-02		7.9E-02		1-
	LA-140	 	0.0E+00		0.00E+00		0.0E+00	 	4

^{*} Includes dose from long-lived progeny
** Includes dose from short-lived progeny

: 1. CAP88PC [Dose Factor	s for Unit C	i Releases	From 331 B	Juilding		
	1			T	<u>_</u>		
	1				<u> </u>		
	fraction of	Offsite		Onsite		·	
	dose from			Worker		Residentia	
X/Q =	parent	2.82E-06		1.49E-05		1.77E-06	
	parent	1000 m N	1				
Location			/ V	200 m S		1000 m E	
Duration	<u> </u>	2000 h/y		2000 h/y		8766 h/y	
Food locally pro	duced	0%		0%		100%	
	<u> </u>	mrem/yr		mrem/yr		mrem/yr	
Radionucl	ide	per Ci rele	ased	per Ci rele	ased	per Ci relea	ased
BA-142		4.5E-05	·	2.36E-04		1.2E-04	
LA-140		2.8E-03		1.46E-02		9.3E-03	
LA-141	1	6.9E-06		3.62E-05		1.8E-05	
LA-142	<u> </u>	3.4E-04		1.80E-03		9.2E-04	
CE-141	 	2.8E-03		1.48E-02		2.7E-02	
CE-143	 	8.2E-04		4.31E-03		2.8E-03	
CE-144	 	6.5E-02		3.43E-01	!	3.7E-01	
PR-143	 	1.4E-03	<u> </u>	7.42E-03		2.5E-02	
	ļ	1	L				
PR-144	<u> </u>	1.1E-05		5.56E-05		2.8E-05	
PR-144M		2.3E-06		1.23E-05		6.3E-06	
ND-147	<u> </u>	1.9E-03	<u> </u>	1.01E-02		2.2E-02	
PM-147	1	6.5E-03		3.42E-02		2.8E-02	
PM-148		3.3E-03		1.75E-02		3.0E-02	
PM-148M		4.7E-02		2.48E-01		2.0E-01	
PM-149		5.6E-04		2.96E-03		3.2E-03	
PM-151		2.2E-04		1.17E-03		6.0E-04	
SM-147		1.2E+01		6.59E+01		3.6E+01	
SM-151	<u> </u>	5.0E-03		2.66E-02		1.8E-02	
SM-153	-	4.3E-04		2.30E-03		2.1E-03	
EU-152	 	1.8E+00		9.46E+00		4.9E+00	
EU-152M		8.6E-05		4.52E-04		2.3E-04	
EU-154		1.4E+00		7.63E+00		4.0E+00	
EU-155		5.7E-02		3.00E-01	• •	1.7E-01	
EU-156		1.1E-02		5.94E-02	·	7.4E-02	
TB-160	<u></u>	3.9E-02		2.05E-01		1.6E-01	
HO-166		6.3E-04		3.31E-03		2.1E-03	
HO-166M		5.9E+00		3.11E+01		1.6E+01	
HF-181		1.3E-02		6.97E-02		7.0E-02	
W-181	 	2.5E-03		1.31E-02		9.5E-03	
W-185		1.3E-04		6.83E-04		1.5E-02	
W-187		3.8E-04		1.99E-03		1.1E-03	
RE-187	-	9.1E-06		4.80E-05	<u> </u>	6.4E-04	
IR-192	ļ	3.4E-02		1.78E-01		1.4E-01	
HG-203		6.2E-03		3.28E-02		4.4E-02	
TL-207		4.0E-07	<u> </u>	2.10E-06		1.1E-06	
TL-208		5.0E-05		2.65E-04	<u> </u>	1.4E-04	
TL-209		1.5E-05		7.70E-05		3.9E-05	
PB-209		1.6E-05		8.52E-05		4.3E-05	
PB-210 **		2.3E+00		1.22E+01		6.3E+01	
PO-210		0.0E+00		0.00E+00		0.0E+00	
PB-211		1.2E-03		6.59E-03		3.4E-03	
PB-212 *	0.93	3.0E-02		1.58E-01		8.1E-02	
PB-214	0.33	1.3E-04		7.01E-04		3.6E-04	
BI-210	ļ	3.3E-02		1.72E-01		9.9E-02	
BI-211	ļ	1.2E-05		6.18E-05	L	3.2E-05	
BI-212		4.9E-03		2.58E-02		1.3E-02	
BI-213		1.8E-04		9.66E-04		4.9E-04	
BI-214		1.5E-04		8.01E-04	L	4.1E-04	
PO-210	T	1.4E+00		7.63E+00	1	2.1E+01	

^{*} Includes dose from long-lived progeny
** Includes dose from short-lived progeny

CAP88PC	Jose Factor	s for Unit C	Releases	From 331 B	ullaing		ļ
			ļ				<u> </u>
	fraction of	Officia		Onsite		ļ	
						Danidantia	<u> </u>
V/O =	dose from	L		Worker		Residentia	
X/Q ≈	parent	2.82E-06	l	1.49E-05		1.77E-06	<u> </u>
Location	· ·	1000 m N\	<u> </u>	200 m S		1000 m E	_
Duration	ļ	2000 h/y		2000 h/y		8766 h/y	
od locally pro	duced	0%		0%		100%	L-
<u> </u>	<u> </u>	mrem/yr	<u> </u>	mrem/yr		mrem/yr	
Radionucl	ide	per Ci rele	ased I	per Ci rele	ased	per Ci rele	ase
PO-218	 	1.2E-06		6.11E-06		3.1E-06	-
RN-219		5.1E-35		2.68E-34		1.4E-34	<u> </u>
RN-220		2.0E-07		1.08E-06		5.5E-07	
RN-222		4.4E-04		2.30E-03		1.2E-03	├
FR-221		1.8E-04		9.70E-04		4.9E-04	├
		3.1E-04		1.63E-03		8.3E-04	
FR-223 RA-223	 	3.1E-04 1.4E+00		7.32E+00		6.6E+00	
RA-223	ļ	5.9E-01		3.10E+00		2.0E+00	<u> </u>
RA-225						4.7E+00	<u> </u>
		7.0E-01		3.69E+00			<u> </u>
RA-226		1.5E+00		7.94E+00		1.7E+01	<u> </u>
RA-228		4.1E-01		2.15E+00		7.9E+00	Ŀ
AC-225		1.0E+00		5.38E+00		3.1E+00	<u> </u>
AC-227		1.1E+02		5.63E+02		3.3E+02	<u> </u>
AC-228		1.4E-02		7.52E-02		3.8E-02	
TH-227	ļ	2.0E+00		1.04E+01		5.5E+00	<u> </u>
TH-228		4.2E+01	·	2.23E+02		1.2E+02	_
TH-229		1.2E+02		6.21E+02		3.2E+02	<u> </u>
TH-230		4.2E+01		2.21E+02		1.2E+02	<u> </u>
TH-231		1.7E-04		9.08E-04		5.4E-04	<u> </u>
TH-232 **	ļ	7.0E+01		3.67E+02		1.9E+02	L
RA-228		0.0E+00		0.00E+00		0.0E+00	L
AC-228		0.0E+00		0.00E+00		0.0E+00	L
TH-228		0.0E+00		0.00E+00		0.0E+00	L
RA-224		0.0E+00		0.00E+00		0.0E+00	Ĺ
RN-220		0.0E+00		0.00E+00		0.0E+00	L
PO-216		0.0E+00		0.00E+00		0.0E+00	L
PB-212		0.0E+00		0.00E+00		0.0E+00	L_
BI-212	<u> </u>	0.0E+00		0.00E+00		0.0E+00	$ldsymbol{f eta}$
TL-208		0.0E+00		0.00E+00		0.0E+00	<u></u>
TH-234		6.1E-03		3.21E-02		9.9E-02	
PA-231		8.0E+01		4.25E+02		2.7E+02	
PA-233		4.5E-03		2.36E-02		3.5E-02	
PA-234		6.1E-04		3.24E-03		1.7E-03	
PA-234M		1.7E-08		8.87E-08		4.5E-08	
U-232		8.0E+01		4.21E+02		2.4E+02	
U-233		2.2E+01		1.17E+02		7.1E+01	
U-234		2.2E+01		1.16E+02		7.0E+01	
U-235		2.1E+01		1.10E+02		6.7E+01	
U-236		2.1E+01		1.10E+02		6.6E+01	
U-237	1	1.1E-03		5.94E-03		1.1E-02	Г
U-238		2.0E+01		1.03E+02	······································	6.2E+01	
U-240		4.1E-04		2.18E-03		1.1E-03	
NP-237		8.2E+01		4.31E+02		2.6E+02	
NP-238		5.9E-03		3.13E-02		1.7E-02	<u> </u>
NP-239		6.7E-04		3.52E-03		3.4E-03	_
NP-240		1.4E-04		7.35E-04		3.7E-04	
NP-240M	 	1.4E-05		7.25E-05		3.7E-05	<u> </u>
PU-236		1.5E+01		7.73E+01		4.4E+01	
PU-238	ļ. ———	5.3E+01		2.79E+02		1.8E+02	

^{*} Includes dose from long-lived progeny
** Includes dose from short-lived progeny

	fraction of		Onsite	
	dose from		Worker	Residential
X/Q =	parent	2.82E-06	1.49E-05	1.77E-06
Location		1000 m NW	200 m S	1000 m E
Duration	<u>L.</u>	2000 h/y	2000 h/y	8766 h/y
Food locally pro	duced	0%	0%	100%
		mrem/yr	mrem/yr	mrem/yr
Radionucl	ide	per Ci released	per Ci released	per Ci released
DI L 020	ļ	E 75 : 04	3.005.00	0.05.00
PU-239	ļ	5.7E+01	3.00E+02	2.0E+02
PU-240	ļ	5.7E+01	2.99E+02	2.0E+02
PU-241		8.6E-01	4.56E+00	3.2E+00
PU-242	ļ	5.4E+01	2.85E+02	1.8E+02
PU-243		4.2E-05	2.22E-04	1.1E-04
PU-244		5.4E+01	2.83E+02	1.8E+02
AM-241		9.0E+01	4.76E+02	2.9E+02
AM-242		9.9E-03	5.25E-02	2.7E-02
AM-242M		8.7E+01	4.59E+02	2.8E+02
AM-243		9.0E+01	4.76E+02	2.9E+02
CM-242		3.0E+00	1.57E+01	9.1E+00
CM-243		6.1E+01	3.20E+02	1.9E+02
CM-244		4.8E+01	2.52E+02	1.5E+02
CM-245	<u> </u>	9.3E+01	4.94E+02	3.0E+02
CM-246		9.2E+01	4.87E+02	2.9E+02
CM-247		8.6E+01	4.52E+02	2.7E+02
CM-248		3.4E+02	1.79E+03	1.1E+03
CF-252		2.6E+01	1.36E+02	7.9E+01
·				

^{*} Includes dose from long-lived progeny
** Includes dose from short-lived progeny

Distribution

No. of <u>Copies</u>				of of opies	
OFFSITE				Pacific Northwest Nation	nal Laboratory
1 DOI	E/Office of Scientific	and Technical		B. P. Atencio	P7-68
Information				M. Y. Ballinger	BSRC
				E. G. Damberg	P7-68
ONSITE				D. L. Edwards (5)	P7-68
	•			T. M. Graham	P7-75
2 DOI	E Richland Operations	Office		S. L. Jones	P7-08
	•			A. S. Ikenberry	P7-79
M. I	E. Burandt	K8-50		K. D. Shields	P7-68
S. D	. Stites	R3-79		M. S. Sula	SEQUIM
				Clearance Office (7)	